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AMENDMENT TO THE CLAIMS

- 1. (Currently Amended): An implantable prosthesis comprising a non-deformable component-comprising a rigid material with pores formed in the rigid material, wherein the rigid material is selected from the group consisting of metals, carbonaceous solids and ceramics, wherein a filler comprising a hydrogel, a structural protein, a bioactive agent, or mixtures thereof, is located within the pores, wherein said rigid porous material with the filler presents a smoother surface for fluid flow than pores without filler,
- 2 (Original): The implantable prosthesis of claim 1 wherein the filler fills the pores.
- 3 (Original): The implantable prosthesis of claim 2 wherein the rigid porous material with the filler presents a smooth surface to flow.
- 4 (Original): The implantable prosthesis of claim 1 wherein the filler partly fills the pores.
- 5 (Original): The implantable prosthesis of claim 1 wherein the filler comprises a hydrogel selected from the group consisting of poly(ethylene glycol), poly(hydroxyethyl methacrylate), partially or fully hydrolyzed poly(vinyl alcohol), poly(vinylpyrrolidone), poly(ethyloxazoline), poly(ethylene oxide)-co-poly(propylene oxide) block copolymers, polyamines, polyacrylamide, hydroxypropylmethacrylate, carboxymethyl cellulose, hydroxyethyl cellulose, methylhydroxypropyl cellulose, polysucrose, hyaluronic acid, alginate, chitosan, dextran, gelatin and mixtures and copolymers thereof.
- 6 (Original): The implantable prosthesis of claim 1 wherein the filler comprises a structural protein.

- 7 (Original): The implantable prosthesis of claim 6 wherein the structural protein is an extracellular matrix protein.
- 8 (Original): The implantable prosthesis of claim 1 wherein the filler comprises a mixture of hydrogel and structural protein.
- 9 (Original): The implantable prosthesis of claim 1 wherein the filler comprises a biologically active agent.
- 10 (Original): The implantable prosthesis of claim 9 wherein the biologically active agent is dispersed within the hydrogel or protein.
- 11 (Original): The implantable prosthesis of claim 9 wherein the biologically active agent is selected from the group consisting of a growth factor, a cell attraction compound, an anticoagulant and combinations thereof.
- 12 (Original): The implantable prosthesis of claim 9 wherein the biologically active agent is VEGF.
- 13 (Original): The implantable prosthesis of claim 9 wherein the bioactive agent is a growth factor.
- 14 (Original): The implantable prosthesis of claim 9 wherein the bloactive agent is a progenitor attraction compound.
- 15 (Original): The implantable prosthesis of claim 9 wherein the bioactive agent is an anticoagulant.

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- 16 (Original): The implantable prosthesis of claim 1 wherein the pores extend through the rigid material.
- 17 (Original): The implantable prosthesis of claim 1 wherein the pores have an interconnecting porosity.
- 18 (Original): The implantable prosthesis of claim 1 wherein a nutrient is also located within the pores.
- 19 (Original): The implantable prosthesis of claim 1 further comprising viable cells.
- 20-21 Canceled.
- 22 (Original): The implantable prosthesis of claim 1 wherein the prosthesis is a mechanical heart valve prosthesis comprising an orifice ring and a rigid occluder attached to the orifice ring.
- 23 (Original): The implantable prosthesis of claim 22 wherein the rigid occluder comprises the rigid material with pores.
- 24-39 Canceled
- 40. (Currently Amended): An implantable medical device comprising a non-deformable component-comprising a rigid material having pores formed in the rigid material and present substantially close to a surface of the rigid material, wherein the rigid material is selected from the group consisting of metals, carbonaceous solids and ceramics, and a filler, wherein said filler comprising a hydrogel, a structural protein, a bioactive agent, or mixtures thereof, is located within the pores to promote cellular attachment and

proliferation.

- 41 (Previously Presented): The medical device of claim 40 wherein said device is for contacting bodily fluids and/or tissue after implantation.
- 42 (Previously Presented): The medical device of claim 40 wherein said filler fills the pores.
- 43 (Previously Presented): The medical device of claim 42 wherein said rigid porous material with the filler presents a smooth surface to flow.
- 44 (Previously Presented): The medical device of claim 40 wherein said bioactive agent is dispersed within the hydrogel or protein.
- 45 (Previously Presented): The medical device of claim 40 wherein the bioactive agent is selected from the group consisting of a growth factor, a cell attraction compound, an anticoagulant and combinations thereof.
- 46 (Previously Presented): The medical device of claim 40 wherein the bioactive agent is VEGF.
- 47 (Previously Presented): The medical device of claim 40 wherein the bioactive agent is a progenitor attraction compound.
- 48 (Previously Presented): The medical device of claim 40 wherein the bioactive agent is an anticoagulant.
- 49 (Currently Amended): An implantable medical device comprising a non-deformable

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component comprising a rigid material having pores substantially extending through the rigid material to form a porous network, wherein the rigid material is selected from the group consisting of metals, carbonaceous solids and ceramics, and a filler, wherein said filler comprising a hydrogel, a structural protein, a bioactive agent, or mixtures thereof, is located within the pores, and said porous network does not provide significant blood flow through the porous material.

- 50 (Previously Presented): The medical device of claim 49 wherein said porous network promotes cellular attachment and proliferation.
- 51 (Previously Presented): The medical device of claim 49 wherein said filler fills the pores.
- 52 (Previously Presented): The medical device of claim 51 wherein said rigid porous material with the filler presents a smooth surface to flow.
- 53 (Previously Presented): The medical device of claim 49 wherein said bioactive agent is dispersed within the hydrogel or protein.
- 54 (Previously Presented): The medical device of claim 49 wherein the bioactive agent is selected from the group consisting of a growth factor, a cell attraction compound, an anticoagulant and combinations thereof.
- 55 (Previously Presented): The medical device of claim 49 wherein the bioactive agent is VEGF.